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(54) Title: METHOD FOR THE SELECTIVE PREPARATION OF 3-OXO-4-AZA-5A-ANDROSTANE COMPOUND

(57) Abstract: This invention relates to a method for selectively preparing the 3-oxo-4-aza-5 α -androstane compound which is used as an intermediate of finasteride by heating 3-oxo-4-aza-5-androstene in a mixture of formic acid and an alkanediol in the presence of zinc.

METHOD FOR THE SELECTIVE PREPARATION OF 3-OXO-4-AZA-5 α -ANDROSTANE COMPOUND ✓

FIELD OF THE INVENTION

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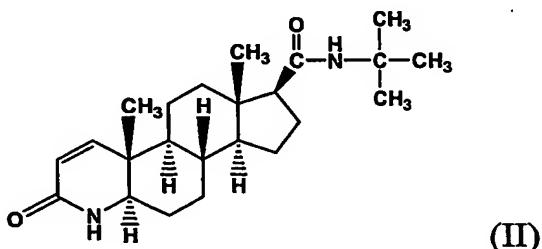
The present invention relates to an improved method for selectively preparing 3-oxo-4-aza-5 α -androstane compound under mild conditions.

DESCRIPTION OF THE PRIOR ART

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Finasteride (17 β -(N-tert-butylcarbamoyl)-5 α -4-aza-androst-1-en-3-on), the compound of formula (II) having an androstane backbone, is effective in treating benign prostatic hypertrophy and androgenetic alopecia:

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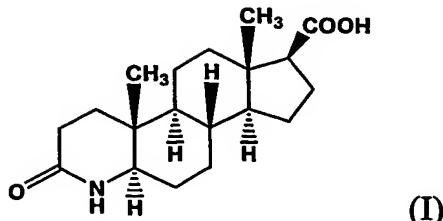
(II)

Benign prostatic hypertrophy and androgenetic alopecia are caused by binding of 5 α -dihydrotestosterone (DHT) derived from testosterone to androgen receptor. The conversion of testosterone into 5 α -dihydrotestosterone is mediated by testosterone 5 α -reductase which is inhibited by finasteride. Such inhibition of 5 α -dihydrotestosterone by finasteride results in rapid recovery of prostate and increased hair growth. Finasteride thus is effective to benign prostatic hypertrophy and good agent for treating androgenic alopecia which exhibits only low, temporary side effects, and it is the only orally administrable among the two hair-growth agents approved by FDA of the United States.

Finasteride can be conventionally prepared by converting the carboxylic group of the 17 β -position of 3-oxo-4-aza-5 α -androstane-17 β -carboxylic acid of formula (I) into a t-butylcarbamoyl group and then carrying out dehydrogenation at the 1,2-positions, or carrying out dehydrogenation at the

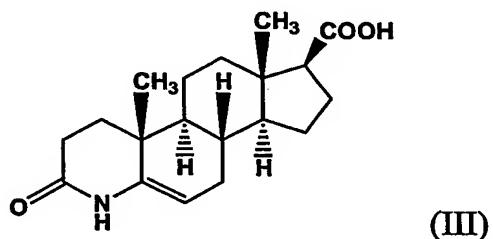
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1,2-positions and then converting the 17β -position carboxylic group into a t-butylcarbamoyl group:



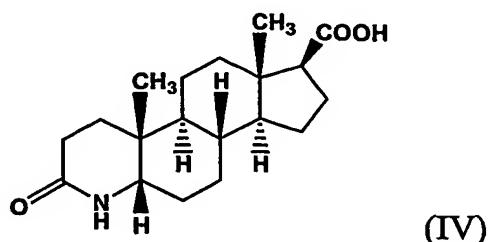
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For example, a process for preparing 3-oxo-4-aza-5 α -androstane- 17β -carboxylic acid of formula (I) is disclosed in U.S. Patent No. 4,760,071 and the *J. Med. Chem.* 29, 2298 (1986), wherein the 3-oxo-4-aza-5-androstene compound of formula (III) is reduced with the 10 hydrogen in the presence of a PtO₂ catalyst under a hydrogen atmosphere of 40psi to produce the compound of formula (I).



15

The above reduction process selectively produces the compound of formula (I) having the 5-hydrogen oriented at 5 α -position, without giving the isomer thereof, the compound of formula (IV) having the 5-hydrogen at the 20 5 β -position. However, this asymmetric reduction process requires the use of explosive hydrogen and an expensive catalyst under high pressure condition.



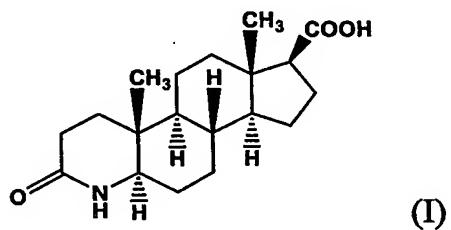
Also disclosed in *J. of Pharmaceutical Sciences.* 63, p 19 (1974) is a method of reducing a steroid compound having a structure similar to the compound of formula (III) to produce a 5 α -compound using formic acid and N-methylformamide. However, this process is conducted under high temperature and high pressure conditions and gives a poor productivity.

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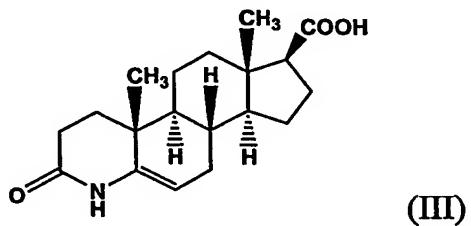
SUMMARY OF THE INVENTION

10 Accordingly, it is a primary object of the present invention to provide an improved method for selectively preparing the compound of formula (I) under mild conditions.

15 In accordance with the present invention, there is provided a method for preparing the compound of formula (I) comprising heating the compound of formula (III) in a mixture of formic acid and an alkanediol in the presence of zinc:



20



BRIEF DESCRIPTION OF THE DRAWINGS

25

The above and other objects and features of the present invention will become apparent from the following description of the invention taken in conjunction with the following accompanying drawings, which respectively

show:

FIG. 1: a high performance liquid chromatography (HPLC) scan of the compound of formula (I) prepared in accordance with the inventive method; and

5 FIG. 2: an HPLC scan of the compound of formula (I) prepared in Comparative Example 1 in the absence of zinc; and

FIG. 3: an HPLC scan of the compound of formula (I) prepared in Comparative Example 2 using formic acid and methylformamide.

10 **DETAILED DESCRIPTION OF THE INVENTION**

The compound of formula (III) used as a starting material of the present invention can be prepared by a conventional method (U.S. Patent No. 4,760,071 and the *J. Med. Chem.* 29, 2298 (1986)).

15 In accordance with the present invention, the compound of formula (I) can be prepared by dissolving the compound of formula (III) in a mixture of formic acid and an alkanediol, adding activated zinc thereto, and heating the resulting mixture.

20 In the inventive method, formic acid may be used in an amount of 3 to 30ml, preferably 5 to 15ml based on 1.0g of the compound of formula (III); and the alkanediol, in an amount of 2 to 20ml, preferably 5 to 10ml, based on 1.0g of the compound of formula (III).

25 The alkanediol which may be used in the present invention includes ethylene glycol, propylene glycol, 1,3-propanediol, 1,2-butanediol, 1,3-butanediol, 1,4-butanediol and 2,3-butandiol, and the like, among which ethylene glycol is preferred.

30 The zinc used in the present invention enhances both the selectivity of the target 5 α -compound and the yield, and also reduces the reaction time. Zinc may be used in 4 to 10 equivalents, preferably, 6 to 8 equivalents, based on a mole of the compound of formula (III), and in the total absence of the isomeric 5 β -byproduct, the target 5 α -compound is produced in a high yield of 80%. When zinc is not used, the target 5 α -compound is produced in a yield of only about 50% together with 10 to 20% of the isomeric 5 β -compound.

35 The reduction in accordance with the present invention may be carried out at a temperature of 80 to 130°C, preferably 100 to 110°C, for 4 to 8 hours.

Thus, in accordance with the simple method of the present invention,

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the target compound of formula (I) can be selectively produced in a high yield under mild conditions.

The present invention will be described in further detail with reference to Examples. However, it should be understood that the present is not
5 restricted by the specific Examples.

Example

Preparation 1: Preparation of 17 β -carboxy-5-oxo-A-nor-3,5-secoandrostan-10 3-onic acid

16g (50mmol) of 3-oxo-4-androstene-17 β -carboxylic acid was dissolved in 240ml of t-butanol, 16g (150mmol) of sodium carbonate dissolved in 40ml of water was added thereto, and then heated to 80 °C. Added dropwise thereto
15 was a solution, which is preheated to 60 °C, of 53.5g (250mmol) of sodium metaperiodate and 4.0g (25mmol) of potassium permanganate dissolved in 300ml of water. The resulting mixture was refluxed for 3 hours and left at room temperature overnight. The inorganic materials were filtered-off through celite, the filtrate was successively washed with water and 250ml of
20 10% sodium hydrogen sulfite, t-butanol was removed under a reduced pressure, and the residue was acidified with concentrated HCl. The acidified residue was then extracted with 320ml of methylene chloride, washed successively with 320ml of 5% sodium hydrogen sulfite and 320ml of brine, and distilled under a reduced pressure, to obtain 14.5g of the title compound (yield: 86%) as
25 a pale yellow solid.

H-NMR(δ , CDCl₃): 0.82(3H, 19-CH₃), 1.16(3H, 18-CH₃), 1.20~2.30 (15H, cyclic-CH), 1.53(2H, 1-CH₂), 2.40(2H, 2-CH₂), 2.50(1H, 17-CH), 11.85(1H, COOH)

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Preparation 2: Preparation of 3-oxo-4-aza-5-androstene-17 β -carboxylic acid (the compound of formula (III))

10g of 17 β -carboxy-5-oxo-A-nor-3,5-secoandrostan-3-onic acid
35 (30mmol) obtained in Preparation 1 was dissolved in 30ml of ethylene glycol, and 75ml of 2.0M ethanolic ammonia solution (150mmol) was added thereto,

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stirred for an hour at 40 to 50°C, and refluxed for 12 hours. The resulting mixture was cooled to room temperature and ethanol was distilled off under a reduced pressure. To the residue was added 150ml of water and the resulting mixture was acidified with 10% HCl to pH 1.5. Precipitates formed were filtered, washed with water, and dried at 45°C, to obtain 6.6g of the title compound (yield: 70%) as a white solid.

H-NMR(δ , DMSO-d₆): 0.57(3H, 19-CH₃), 0.91(3H, 18-CH₃), 0.95~2.30 (18H, cyclic-CH), 4.76(1H, 6-CH), 9.17(1H, NH), 11.85(1H, COOH)

10

Example 1: 3-oxo-4-aza-5 α -androstane-17 β -carboxylic acid (the compound of formula (I) - 1)

3.2g (10mmol) of 3-oxo-4-aza-5-androstene-17 β -carboxylic acid obtained in Preparation 2 was dissolved in a mixture of 45ml of formic acid and 15ml of ethylene glycol, and 2.6g (80mmol) of activated zinc was added thereto. The mixture was reacted for 8 hours at 100 to 105°C and cooled to room temperature. The suspended solid was removed by filtration, and the solvent in the filtrate was removed under a reduced pressure. 13ml of N-methylformamide was added to the residue, and the resulting mixture was stirred for 30 minutes in an ice bath. Precipitates formed were then filtered and dried at 45°C, to obtain 2.6g of the title compound (yield: 81%) as a white solid.

The product thus obtained was analyzed by HPLC and the result is shown in FIG. 1. As can be seen in FIG. 1, only the target 5 α -compound (retention time: 11.996) is detected, the isomeric 5 β -compound being not detectable.

H-NMR(δ , DMSO-d₆): 0.56(3H, 19-CH₃), 0.72(3H, 18-CH₃), 0.80~1.30 (8H, cyclic-CH), 1.40~1.70(7H, cyclic-CH), 1.87(2H, 16-CH), 2.10(2H, 2-CH₂), 2.30(1H, 17-CH), 3.0(1H, 5-CH), 7.15(1H, NH), 11.85(1H, COOH)

Example 2: 3-oxo-4-aza-5 α -androstane-17 β -carboxylic acid (the compound of formula (I) - 2)

35

3.2g (10mmol) of 3-oxo-4-aza-5-androstene-17 β -carboxylic acid

obtained in Preparation 2 was dissolved in a mixture of 16ml of formic acid and 32ml of ethylene glycol, and 2.6g (80mmol) of activated zinc was added thereto. The mixture was reacted for 8 hours at 110 to 120°C, and cooled to room temperature. The suspended solid was removed by filtration, formic
5 acid was removed under a reduced pressure. The residue was dissolved in 300ml of chloroform and washed successively with 150ml portions of 5% aqueous sodium carbonate solution (x2) and 150ml portions of water (x3). The chloroform layer was separated, then dried, filtered and the solvent was removed under a reduced pressure. 13ml of N-methylformamide was
10 added to the residue and stirred for 30 minutes in an ice bath. Precipitates formed were then filtered and dried at 45°C, to obtain 2.7g of the title compound (yield: 83%) as a white solid.

The product thus obtained was analyzed by HPLC and the result showed that only the 5 α -compound (retention time: 11.996) was produced.
15 H-NMR data was the same as in Example 1.

Comparative example 1: Preparation of 3-oxo-4-aza-5 α -androstane-17 β -carboxylic acid (the compound of formula (I)) in the absence of zinc

20 3.2g (10mmol) of 3-oxo-4-aza-5-androstene-17 β -carboxylic acid obtained in Preparation 2 was dissolved in a mixture of 45ml of formic acid and 15ml of ethylene glycol, and reacted for 8 hours at 100 to 105°C. The reaction mixture was cooled to room temperature, the residual solid was remove by filtration and the solvent was distilled off under a reduced
25 pressure. 13ml of N-methylformamide was added to the resulting residue and stirred for 30 minutes in an ice bath. Precipitates formed were then filtered and dried at 45°C, to obtain 1.7g of the title compound (yield: 53%) as a white solid.

The product thus obtained was analyzed by HPLC and the result is
30 shown in FIG. 2, wherein the ares of 5 β -compound peak (retention time: 12.956) is 15% relative to the area of the 5 α -compound peak (retention time: 12.187) of 85%. That is, a large amount of the undesired 5 β -compound is produced.

carboxylic acid (the compound of formula (I)) using a mixture of formic acid and N-methylformamide

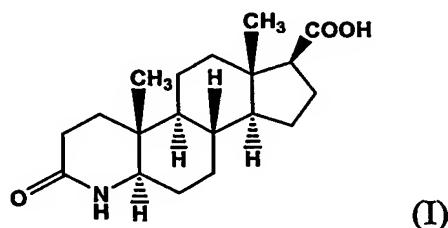
3.2g (10mmol) of 3-oxo-4-aza-5-androstene-17 β -carboxylic acid obtained in Preparation 2 was dissolved in a mixture of 45ml of formic acid and 15ml of N-methylformamide, and reacted for 8 hours at 100 to 105 °C. The reaction mixture was cooled to room temperature, the residual solid was filtered off, formic acid was removed under a reduced pressure, and the remaining solution was stirred for 30 minutes in an ice bath. Precipitates formed were then filtered and dried at 45 °C, to obtain 1.9g of the target compound (yield: 59%) as a white solid.

The product thus obtained was analyzed by HPLC and the result is shown in FIG. 3, wherein the area of the 5 β -compound peak (retention time: 12.770) is 35% relative to the 5 α -compound peak (retention time: 12.046) of 65%. That is, a large amount of the undesired 5 β -compound is produced.

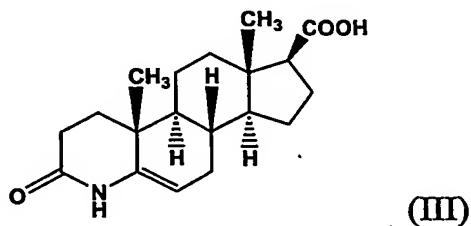
While the invention has been described with respect to the specific embodiments, it should be recognized that various modifications and changes may be made by those skilled in the art to the invention which also fall within the scope of the invention as defined as the appended claims.

WHAT IS CLAIMED IS:

1. A method for preparing the compound of formula (I) comprising heating the compound of formula (III) in a mixture of formic acid and an alkanediol in the presence of zinc:



1.0



2. The method of claim 1, wherein formic acid is used in an amount of 3 to 30ml based on 1.0g of the compound of formula (III).

15

- 3 The method of claim 1, wherein the alkanediol is selected from the group consisting of ethylene glycol, propylene glycol, 1,3-propanediol, 1,2-butanediol, 1,3-butanediol, 1,4-butanediol and 2,3-butandiol.

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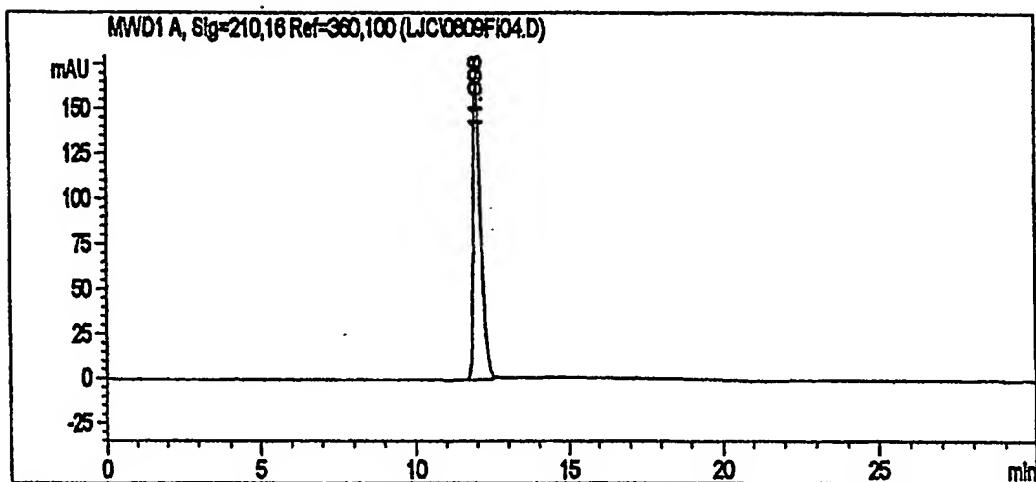
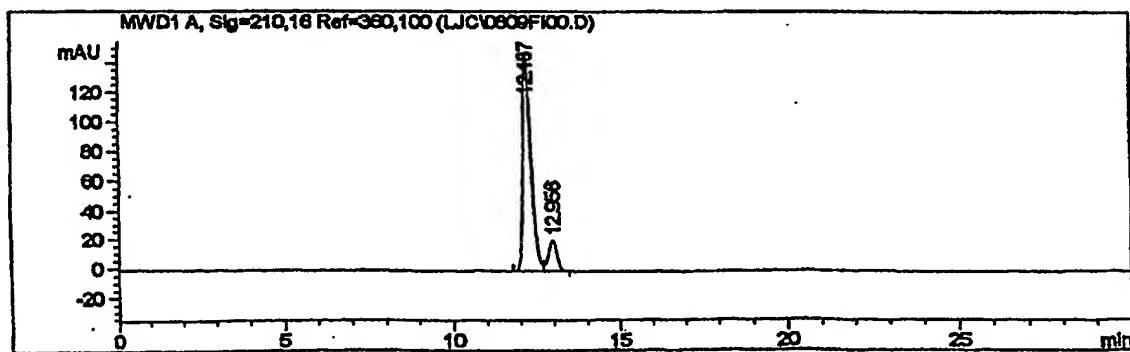
4. The method of claim 1, wherein the alkanediol is used in an amount of 2 to 20ml based on 1.0g of the compound of formula (III).

5. The method of claim 1, wherein zinc is used in 4 to 10 equivalents based on a mole of the compound of formula (III).

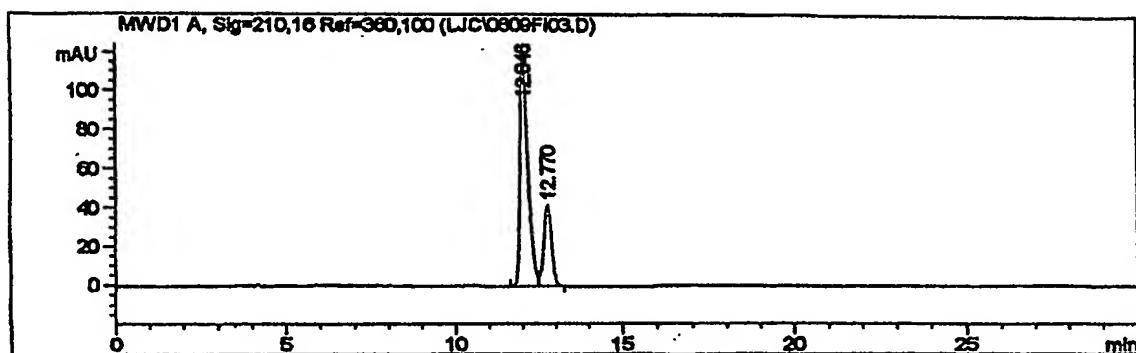
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6. The method of claim 1, wherein the heating is carried out for 4 to 8 hours at a temperature in the range of 80 to 130°C.

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FIG. 1**FIG. 2**

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FIG. 3

INTERNATIONAL SEARCH REPORT

International application No.
PCT/KR03/01629

A. CLASSIFICATION OF SUBJECT MATTER

IPC7 C07D 221/18

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

C07D 221/18

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)
STN online, CA on CD, KIPASS

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 02/46207 A2 (Glaxo group limited) 13 June 2002 (2002-06-13) page7, line 7-30 example 1-2	1-6
A	US 5804576 A (Research Corporation Technologies, Inc.) 8 September 1998 (1998-09-08) scheme 1 section 37, line 49-50	1-6
A	US 4451405 A (Glaxo group limited) 29 May 1984 (1984-05-29) section 5, line 63-65	1-6
A	John F. Templeton, Samar Majid, and Angelina Marr, 'Zinc-Acetic Acid Reduction of the Steroid 4-En-3-one: Novel Conversion of the 4-En-3-one into the 2-En-4-one via a Vinyl Chloride, J. Chem. Soc. Perkin Trans. 1, 1990, Vol.9, p.2581-2584 page 2581 left column line 8-18; claim 1 page 2582 Experimental section 2a; claim 1	1
A	Peng Xia et al., 'Synthesis of N-substituted 3-Oxo-17beta-carboxamide-4-aza-5alpha-androstanes and the Tautomerism of 3-Oxo-4-aza-5-androstenes', Heterocycles, 1998, Vol.47(2), p.703-716 page 704, line 8-20; claim 1 page 705, line 12-page 706, line 2; claim 1	1

Further documents are listed in the continuation of Box C.

See patent family annex.

* Special categories of cited documents:	
"A"	document defining the general state of the art which is not considered to be of particular relevance
"E"	earlier application or patent but published on or after the international filing date
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Date of the actual completion of the international search 25 NOVEMBER 2003 (25.11.2003)	Date of mailing of the international search report 26 NOVEMBER 2003 (26.11.2003)
Name and mailing address of the ISA/KR  Korean Intellectual Property Office 920 Dunsan-dong, Seo-gu, Daejeon 302-701, Republic of Korea Facsimile No. 82-42-472-7140	Authorized officer LEE, HYUN SONG Telephone No. 82-42-481-5606



INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No.

PCT/KR03/01629

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